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(54) **Aqueous preparation of azulene sodium sulfonate.**

(57) Aqueous solution preparations which comprise azulene sodium sulfonate as an active ingredient and 20 w/v% or more of a polyhydric alcohol.

The active ingredient, azulene sodium sulfonate, is maintained stably in the preparation for a prolonged period of time. Moreover, the preparation is highly safe and suited for the easy and hygienic use for treating oral cavity and/or throat.

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BACKGROUND OF THE INVENTION

Field of the Invention:

5 The present invention relates to stable aqueous solution preparations containing azulene sodium sulfonate as an active ingredient, which are suited for use as a preventive or therapeutic medicine for inflammatory diseases in digestive organs, oral cavity and throat, otolaryngology, ophthalmology, etc., and which can be used easily and hygienically.

10 Discussion of the Background:

Azulene sodium sulfonate is found in an extract of the plant camomile (*Matricaria Chamomilla* L.) which belongs to the Asteraceae originated in the South and East Europe. As a folk medicine, it has long been used to cure a variety of inflammatory diseases. Currently, it is widely used as an anti-inflammatory agent to be incorporated in medicines for treating gastritis and gastric ulcer, mouthwashes, oral preparations, eye drops and the like.

15 Azulene sodium sulfonate is generally formed into solids such as tablets, lozenges, granule, powder, and the like, with an exception of eye drops. Drugs for internal use are orally administered as they are, while mouthwashes are prepared by dissolving the solid drug in water just before dosing. This is primarily due to the fact that azulene sodium sulfonate is unstable and easily degraded in the aqueous solution as time passes and under diffused light.

20 Eye drops, which are the only preparation marketed in liquids, usually contain adjuvants such as thiosulfate (antioxidizing agent) and EDTA (chelating agent), and are provided in expensive air-tight containers. Some of the adjuvants cannot be used as ingredients of drugs for oral administration. Moreover, the stability of eye drop preparations is not sufficiently good.

25 Reflecting the increased number of patients suffering from oral diseases typified by stomatitis and sore throat, which may possibly be due to the overpopulation and environmental pollution, there has recently been a growing demand for OTC (over-the-counter) drugs which are easily applied to the oral cavity/throat and are free from adverse side effects. As a solution, lozenge or dissolving-on-use type preparations containing azulene sodium sulfonate are considered. However, they are not necessarily satisfactory in that the active ingredient, azulene sodium sulfonate, will not successfully reach the affected area via lozenge, while dissolving-on-use type will require water and a glass, which makes the whole process troublesome and non-hygienic.

30 Accordingly, there is still a demand for an aqueous solution preparation of azulene sodium sulfonate suited for the internal administration or the oral cavity/throat application. So far before the present invention, however, it has been considered difficult to industrially provide such an aqueous preparation containing azulene sodium sulfonate as an active ingredient, because the sulfonate is unstable in an aqueous solution, and hence, aqueous solution preparations of azulene sodium sulfonate have never existed.

40 SUMMARY OF THE INVENTION

Accordingly, it is one object of the present invention is to provide a stable aqueous solution preparation containing azulene sodium sulfonate as an active ingredient.

45 It is another object of the present invention to provide an aqueous solution preparation which comprises azulene sodium sulfonate as an active ingredient and 20 w/v% or more of a polyhydric alcohol.

These and other objects, features and advantages of the present invention will become apparent from the following description.

DETAILED DESCRIPTION OF THE INVENTION AND PREFERRED EMBODIMENTS

50 The present inventors have conducted extensive studies and have found that when an aqueous solution preparation of azulene sodium sulfonate contains a polyhydric alcohol as a solubilizer, the stability of azulene sodium sulfonate is maintained for an unexpectedly prolonged period of time.

55 The polyhydric alcohol is known to be a highly safe substance. Examples of the polyhydric alcohol useful in the present invention include glycerin, glycols and sugaralcohols. As for glycols, mention may be given to propylene glycols and the like. As for sugaralcohols, mention may be given to sorbitol, xylitol and the like. Among them, glycerin is particularly preferred from the viewpoint of the excellent stabilizing ability for azulene sodium sulfonate.

These polyhydric alcohols are contained in the aqueous solution preparation according to the present invention in amounts of 20 w/v% (hereinafter simply referred to as %) or more, preferably from 20 to 90%, and more preferably from 30 to 80%. Amounts less than 20% are not preferred because azulene sodium sulfonate cannot be maintained stably for a prolonged period of time.

The present aqueous solution preparation can contain optional ingredients so long as they do not impede the effects of the present invention. Examples of such optional ingredients include flavors, buffers, preservatives and stabilizers.

The aqueous solution preparation according to the present invention can be prepared by conventional methods, for instance, by blending a mixture of azulene sodium sulfonate and a polyhydric alcohol with a buffer, preservative, sterile distilled water, etc. Although no particular limitation is imposed on the concentration of azulene sodium sulfonate, it is preferred that the concentration be from 0.02 - 0.06%. Moreover, the pH of the aqueous solution preparation according to the present invention is preferably adjusted to fall within the range from 7 - 9.

The thus-obtained aqueous solution preparation of the present invention is useful as medicines for internal administration, oral cavity/throat application or nasal drops. In case of the oral cavity/throat application, it is preferred that the preparation be placed in a handy container which is so designed that one can apply the right amount of dose by directly jet-spraying onto the affected part. This combination of the aqueous solution preparation of azulene sodium sulfonate and the handy container for jet-spraying is especially advantageous in the treatment of inflammation of the oral cavity and pharyngitis, because water is no more required.

EXAMPLES

The present invention will now be described more specifically by way of examples, which however, should not be construed as limiting the invention thereto.

Example 1.

(1) Aqueous 0.02% solutions of azulene sodium sulfonate were added with 80% polyhydric alcohols (glycerin, sorbitol, xylitol and propyleneglycol, separately) to prepare aqueous test solutions. Each of the test solutions was charged in a 15 ml polyethylene container, and stored in a thermo-hydrostat under the conditions of $40 \pm 1^\circ\text{C}$ and $75 \pm 5\% \text{RH}$ (light shielded). The content of azulene sodium sulfonate was measured by an auto-recording spectrophotometer (manufactured by Shimazu, UV-240) at the wavelength of 570 nm as time elapsed. The residual rate was calculated counting the absorbance measured just preceding the storage as 100%. The results are shown in Table 1. The data show that the addition of polyhydric alcohol gives significantly higher stabilizing effect when compared to the absence of polyhydric alcohol. Specifically, glycerin exhibited outstanding and remarkable stabilizing effect, indicating that azulene sodium sulfonate should be stable over 3 years when stored at the normal ambient temperature.

Table 1

	Just before storage	After 2 wks	After 4 wks	After 6 wks	After 8 wks
Glycerin	100.0	100.0	100.0	99.2	98.7
Sorbitol	100.0	93.7	86.7	82.6	80.0
Xylitol	100.0	95.9	90.0	86.3	82.2
Propylene glycol	100.0	96.8	90.6	85.7	81.0
(non addition)	100.0	92.6	85.2	75.9	66.7

(2) Aqueous 0.02% azulene sodium sulfonate solutions were added with 0%, 20%, 40%, 60% or 80% of glycerin, respectively, and charged in transparent glass vials. Stability against the light was investigated by following the similar procedure as described in (1) above. The residual rate was calculated counting the absorbance just before the storage as 100%. The results are shown in Table 2. The data show that the stabilizing effect is more significant as the concentration of glycerin increases. When 80% of glycerin was added, the residual rate after 4 hour storage was 94.2% indicating very high stability, whereas

absence of glycerin and 20% of glycerin gave the residual rates of only 54.8% and 72.4%, respectively. Thus, the stabilizing activity of glycerin was statistically meaningful.

Table 2

	Exposure to the sunlight				
	0 hr	1 hr	2 hrs	3 hrs	4 hrs
Glycerin 0%	100.0	81.2	72.1	64.5	54.8
Glycerin 20%	100.0	91.3	84.3	78.9	72.4
Glycerin 40%	100.0	94.8	91.4	87.4	83.0
Glycerin 60%	100.0	97.5	96.5	93.7	90.9
Glycerin 80%	100.0	97.7	97.0	95.7	94.2

(3) Aqueous 0.02% azulene sodium sulfonate solutions added with 0%, 20%, 40%, 60% and 80% of glycerin, respectively, were stored in a thermo-hydrostat under the same severe conditions as described in (1) to study changes in the residual rate as time elapses over 12 weeks. As a result, higher concentration of glycerin provided more significant stabilizing effect as shown in Table 3. Practically, it is advisable that azulene sodium sulfonate should be maintained at a residual rate of over 90% after 12 weeks of storage (equivalent to 1.5 year storage at room temperature). From this standard, it is determined that the minimal concentration of glycerin to be added is 20% in a practical use.

Table 3

	Just before Storage	after 4w storage	after 8w storage	after 12w storage
Glycerin 0%	100.0	85.2	66.7	44.8
Glycerin 20%	100.0	98.2	95.0	91.4
Glycerin 40%	100.0	99.2	97.0	94.0
Glycerin 60%	100.0	100.0	98.4	97.2
Glycerin 80%	100.0	100.0	98.7	97.4
Storage conditions: $40 \pm 1^\circ\text{C}$, $75 \pm 5\% \text{RH}$ (light shielded)				

Example 2.

Formulation:

Azulene sodium sulfonate	0.02 g
Glycerin	25
Sodium hydrogencarbonate	0.2
Sodium sorbate	0.1
Flavor	suitable amount

The above ingredients were mixed and added with sterile distilled water to dissolve and to make the total quantity of 100 ml. The pH of the solution was adjusted to approximately 8.2. Subsequent to the aseptic filtration, the solution was charged in an air-tight container.

Example 3.

Formulation:

Azulene sodium sulfonate	0.02 g
Glycerin	35
Sodium citrate	0.2 g
Ethanol (95%)	15 ml
L-menthol	0.01 ml

The above ingredients were mixed and added with sterile distilled water to dissolve and to make the total quantity of 100 ml. The pH of the solution was adjusted to approximately 8.4. Subsequent to the aseptic filtration, the solution was charged in an air-tight container.

Example 4.

Formulation:

Azulene sodium sulfonate	0.04 g
Glycerin	80
Boric Acid	1
Borax	0.4
Paraoxypropyl benzoate	0.07
Paraoxymethyl benzoate	0.13

The above ingredients were mixed and added with sterile distilled water to dissolve and to make the total quantity of 100 ml. The pH of the solution was adjusted to approximately 7.8. Subsequent to the aseptic filtration, the solution was charged in an air-tight container.

Example 5.

Formulation:

Azulene sodium sulfonate	0.02 g
Sorbitol	75
Boric acid	1
Borax	0.4
Ethylenediamine tetraacetate	0.05
Sodium thiosulfate	0.2

The above ingredients were mixed and added with sterile distilled water to dissolve and to make the total quantity of 100 ml. The pH of the solution was adjusted to approximately 8.3. Subsequent to the aseptic filtration, the solution was charged in a nasal spray container.

As described herein above in this specification, the present invention provides an aqueous solution preparation containing azulene sodium sulfonate as an active ingredient, which is stable over a prolonged period of time, is highly safe, and is suited for handy and hygienic use in the treatment of the oral cavity and throat.

The features disclosed in the foregoing description, in the claims and/or in the accompanying drawings may, both, separately and in any combination thereof, be material for realizing the invention in diverse forms thereof.

Claims

1. An aqueous solution preparation which comprises azulene sodium sulfonate as an active ingredient and 20 w/v% or more of a polyhydric alcohol.
- 5 2. The preparation as defined in Claim 1, wherein the polyhydric alcohol is selected from the group consisting of glycerin, glycols and sugaralcohols.
3. The preparation as defined in Claim 1, wherein the polyhydric alcohol is glycerin.
- 10 4. The preparation as defined in Claim 1, wherein the amount of the azulene sodium sulfonate is from 0.02 to 0.06 w/v% and the amount of the polyhydric alcohol is from 20 to 90 w/v%.
- 15 5. The preparation as defined in Claim 4, further comprising optional ingredients selected from the group consisting of flavors, buffers, preservatives, stabilizers and water.
6. A method for the treatment of oral cavity or throat, which comprises applying the preparation of Claim 1 to the affected part.



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PARTIAL EUROPEAN SEARCH REPORT

Application Number

which under Rule 45 of the European Patent Convention EP 93 11 1145 shall be considered, for the purposes of subsequent proceedings, as the European search report

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.5)
P,X	DATABASE WPI Section Ch, Week 9248, Derwent Publications Ltd., London, GB; Class B05, AN 92-395343 & JP-A-4 295 419 (NIPPON SHINYAKU CO. LTD.) 20 October 1992 * abstract *	1-5	A61K31/015
Y	US-A-4 525 343 (H. RAAF) * example 6 *	1-5	-
Y	DATABASE WPI Section Ch, Week 8601, Derwent Publications Ltd., London, GB; Class B05, AN 86-003026 & JP-A-60 228 423 (NITTO ELECTRIC IND. KK.) 13 November 1985 * abstract *	1-5	-
			TECHNICAL FIELDS SEARCHED (Int.Cl.5)
			A61K
INCOMPLETE SEARCH			
<p>The Search Division considers that the present European patent application does not comply with the provisions of the European Patent Convention to such an extent that it is not possible to carry out a meaningful search into the state of the art on the basis of some of the claims</p> <p>Claims searched completely : Claims searched incompletely : Claims not searched : Reason for the limitation of the search:</p> <p>see sheet C</p>			
Place of search MUNICH		Date of completion of the search 25 January 1994	Examiner Foerster, W
CATEGORY OF CITED DOCUMENTS			
<p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			



EP 93 11 1145 -C-

INCOMPLETE SEARCH

Claims searched completely : 1-5
Claims not searched : 6

Reason : Art 52(4).